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(54) **Process for the drying of a hydrocarbon stream**

(57) The invention concerns a process for the continuous drying of a hydrocarbon stream and comprises contacting the hydrocarbon stream with an ionic, liquid drying agent of a salt of a fluorinated sulphonic acid.

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triflate in a counter flow contact column 2 at a temperature of 20°C in a continuous operation. The ionic liquid was charged through line 10 at a feed rate of 109 g/h to the top of the column and withdrawn at the bottom. The hydrocarbon stream was charged at a feed rate of 3383 g/h to the bottom of the contact column and withdrawn from the top. The contact column was 0.5m high with an ID=20 mm and packed with 200 ml 4 mm glass helices.

[0020] The wetted ionic liquid withdrawn from the contact column 2 was charged to the top of stripping column 6 in which the ionic liquid was dried by stripping in counter flow with a stream 8 of superheated heptane (150°C) before being withdrawn at the bottom of the stripping column and returned to the top of the contact column. The wet ionic liquid was charged to the top at a feed rate of 109 kg/h at ambient temperature and the dried ionic liquid was withdrawn from the bottom of the stripping column and returned to the hydrocarbon contact column.

[0021] Heptane vapour was fed to the stripping column at 1140 g/h. The stripping column was 0.4 m high with ID = 30 mm and filled with 300 ml 4 x 4 mm stainless steel Rashig rings.

[0022] The results are given in Tables 1 and 2 below.

Table 1

| Contact Column<br>(Hydrocarbon drying)                  |      |
|---|------|
| Hydrocarbon flow ( feed, g/h)                           | 3383 |
| Ionic liquid flow, (g/h)                                | 109  |
| Before treatment<br>(Wet hydrocarbon feed, ppm water)   | 80   |
| After treatment<br>(Dry hydrocarbon product, ppm water) | 5.5  |
| Hydrocarbon feed temperature (°C)                       | 20   |
| Ionic liquid feed temperature (°C)                      | 20   |
| Hydrocarbon / Ionic liquid rate                         | 31   |

[0023] The results show that the water content in the hydrocarbon feed is reduced from 80 ppm to 5.5 ppm.

Table 2

| Stripping Column<br>(Ionic liquid drying)          |         |
|--|---------|
| Ionic liquid flow, (g/h)                           | 109     |
| Heptane flow, (g/h)                                | 1140    |
| Before treatment<br>(Wet ionic liquid, ppm water)  | 2632    |
| After treatment<br>(Dried ionic liquid, ppm water) | 198     |
| Ionic liquid feed temperature (°C)                 | Ambient |

Table 2 (continued)

| Stripping Column<br>(Ionic liquid drying) |    |
|---|----|
| Heptane feed temperature (°C)             | 98 |

### Example 2

[0024] This experiment was performed in the same equipment as used for Example 1. However, the flows were changed, as illustrated in Tables 3 and 4 below, to give a hydrocarbon flow/ionic liquid flow ratio of 4 in the contact column. The actual flows are shown in the Tables.

Table 3

| Contact Column<br>(Hydrocarbon drying)                  |      |
|---|------|
| Hydrocarbon flow (feed, g/h)                            | 1609 |
| Ionic liquid flow, (g/h)                                | 409  |
| Before treatment<br>(Wet hydrocarbon feed, ppm water)   | 76   |
| After treatment<br>(Dry hydrocarbon product, ppm water) | 3.5  |
| Hydrocarbon feed temperature (°C)                       | 20   |
| Ionic liquid feed temperature (°C)                      | 20   |
| Hydrocarbon / Ionic liquid rate                         | 4    |

[0025] The results indicated in Table 3 show that the water content in the hydrocarbon stream was reduced from 76 ppm to 3.5 ppm.

Table 4

| Stripping Column<br>(Ionic liquid drying)         |         |
|---|---------|
| Ionic liquid flow, (g/h)                          | 409     |
| Heptane flow, (g/h)                               | 1140    |
| Before treatment<br>(Wet ionic liquid, ppm water) | 217     |
| After treatment<br>(Dry ionic liquid, ppm water)  | 88      |
| Ionic liquid feed temperature (°C)                | Ambient |
| Heptane feed temperature (°C)                     | 98      |

### Claims

1. Process for the continuous drying of a hydrocarbon stream comprising contacting the hydrocarbon stream with an ionic, liquid drying agent of a salt of a fluorinated sulphonic acid.

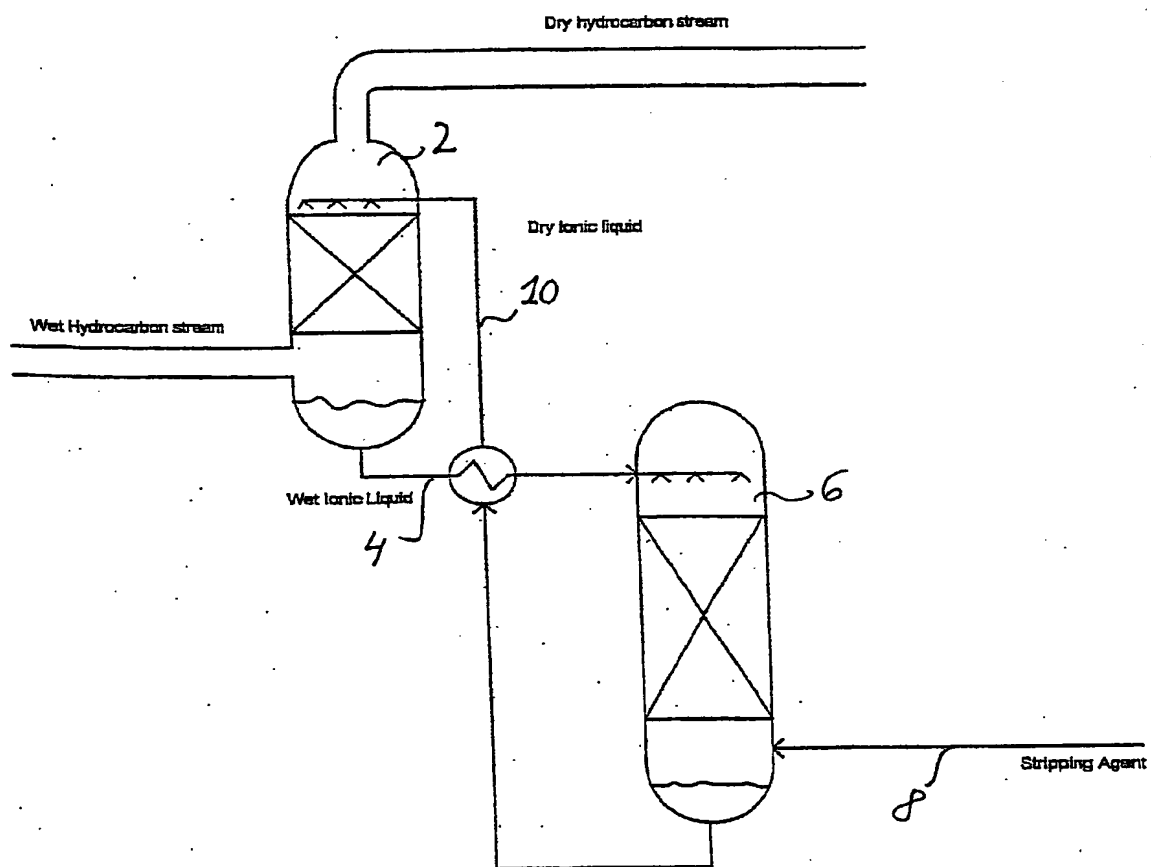


Fig. 1

**ANNEX TO THE EUROPEAN SEARCH REPORT  
ON EUROPEAN PATENT APPLICATION NO.**

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| Patent document<br>cited in search report |   | Publication<br>date | Patent family<br>member(s) | Publication<br>date |
|---|---|---------------------|----------------------------|---------------------|
| DE 2607258                                | A | 01-09-1977          | DE 2607258 A1              | 01-09-1977          |
| US 5725637                                | A | 10-03-1998          | AU 5005397 A               | 22-05-1998          |
|   |   |                     | WO 9818542 A1              | 07-05-1998          |
|   |   |                     | US 5725636 A               | 10-03-1998          |
|   |   |                     | US 5462584 A               | 31-10-1995          |
| GB 1426401                                | A | 25-02-1976          | NONE                       |                     |

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